



LABORATORY REPORT

NAME : MR.SI0067 REFERRED BY : SELF VISIT NO : VAMP26148029
AGE : 40Y 0M 0D ZERO TARIFF CLIENT CODE COLLECTED ON : 21-04-2026 10:00
GENDER : Male LAB MR# : AAMP01479308 RECEIVED ON : 21-04-2026 18:11
OP / IP / DG # : APPROVED ON : 22-04-2026 13:42
REPORT STATUS : Final Report



Test Name	Result	Biological Ref. Interval	Unit
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SEROLOGY AND IMMUNOLOGY

Dengue IgG antibody (Serum)

Dengue IgG antibody 6.30 <18:Negative UNITS
ELISA 18-22:Equivocal
>22:Positive

Interpretation:

RESULT	INTERPRETATION
NEGATIVE (<18)	No detectable IgG antibody, presumptive evidence that the patient does not have secondary Dengue infection.
EQUIVOCAL (18-22)	Repeat sample after 1 week
POSITIVE (>22)	IgG antibody detected indicating presumptive evidence that the patient has been recently exposed to / or currently infected with Dengue virus.

Dengue is considered to be the most important arthropod borne viral disease due to the human morbidity and mortality it causes. The disease may be subclinical, self limiting, febrile or may progress to a severe form of Dengue hemorrhagic fever or Dengue shock syndrome. The clinical diagnosis must be interpreted with clinical signs and symptoms of the patient. The positive predictive value depends on the likelihood of the virus being present. Secondary dengue virus infection causes complications and is characterized by high IgG levels detectable as early as 2 days following the onset of infection, which may be accompanied by elevated IgM levels. The accuracy of the assay is dependent upon the timing following the infection when the sample is collected. Although this assay can detect samples as early as three days following infection, the peak performance of the assay is achieved when samples are taken between 6–15 days following onset of illness.

Dr. G. Amitha
MBBS, MD (MICROBIOLOGY)
Consultant Microbiologist

Disclaimer:

- All results released pertain to the specimen as received by the lab for testing and under the assumption that the patient indicated or identified on the bill/test requisition form is the owner of the specimen.
- Clinical details and consent forms, especially in Genetic testing, histopathology, as well as wherever applicable, are mandatory to be accompanied with the test requisition form. The non-availability of such information may lead to delay in reporting as well as misinterpretation of test results. The lab will not be responsible for any such delays or misinterpretations thereof.
- Test results are dependent on the quality of the sample received by the lab. In case the samples are preprocessed elsewhere (e.g., paraffin blocks), results may be compromised.
- Tests are performed as per the schedule given in the test listing and in any unforeseen circumstances, report delivery may be affected.





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<p>5. Test results may show inter-laboratory as well as intra-laboratory variations as per the acceptable norms.</p> <p>6. Genetic reports as well as reports of other tests should be correlated with clinical details and other available test reports by a qualified medical practitioner. Genetic counselling is advised in genetic test reports by a qualified genetic counsellor, medical practitioner or both.</p> <p>7. Samples will be discarded post processing after a specified period as per the laboratory's retention policy. Kindly get in touch with the lab for more information.</p> <p>8. If accidental damage, loss, or destruction of the specimen is not attributable to any direct or negligent act or omission on the part of Ampath Labs or its employees, Ampath shall in no event be liable. Ampath lab's liability for a lack of services, or other mistakes and omissions, shall be restricted to the amount of the patient's payment for the pertinent laboratory services.</p>			

